Current Biology Dispatches



There are, however, alternative possibilities. Current models of optomotor responses and fixation behaviour in flies include motion-sensitive sensory pathways that provide feedback to turn the animals^{4,9,10}. Interestingly, tethered flies walking on a track ball take a stable heading direction with respect to a stripe projected to the animals in closed loop. This behaviour involves central circuits that calculate heading error by comparing sensory information with an internally represented goal orientation. In this model, however, the error signal triggers also directed turning manoeuvres¹¹. These behavioural strategies are different to the ones proposed in the current study. Thus, future work should show whether these differences are due to the different

behavioural paradigms employed or whether the various models can perhaps be aligned. *Drosophila* is a tractable experimental system that provides the opportunity to further scrutinize these various concepts.

It would also be interesting to test how flies perform under more natural conditions where they walk on rough terrain that causes a high degree of movement noise. It is conceivable that posture stabilization would get prioritized in such situations. This would imply that the fly brain is able to actively regulate the relative contributions of the gaze- and posture-stabilizer. This could be of ethological relevance for flies walking in the unpredictable terrains of natural habitats. In accordance with this view, Cruz et al.² found that visual feedback is not effective during saccades, suggesting that indeed motor context is crucially controlling the relative roles of proprioceptive versus visual feedback.

REFERENCES

- Matthis, J.S., Yates, J.L., and Hayhoe, M.M. (2018). Gaze and the control of foot placement when walking in natural terrain. Curr. Biol. 28, 1224–1233.
- Cruz, T.L., Pérez, S.M., and Chiappe, M.E. (2021). Fast tuning of posture control by visual feedback underlies gaze stabilization in walking Drosophila. Curr. Biol. 31, 4596–4607.

- CellPress
- Borst, A. (2014). Fly visual course control: behaviour, algorithms and circuits. Nat. Rev. Neurosci. 15, 590–599.
- 4. Bahl, A., Ammer, G., Schilling, T., and Borst, A. (2013). Object tracking in motion-blind flies. Nat. Neurosci. *16*, 730–738.
- Costa, A.C., Ahamed, T., and Stephens, G.J. (2019). Adaptive, locally linear models of complex dynamics. Proc. Natl. Acad. Sci. USA *116*, 1501–1510.
- Bizzi, E., Kalil, R.E., and Tagliasco, V. (1971). Eye-head coordination in monkeys: evidence for centrally patterned organization. Science 173, 452–454.
- DeAngelis, B.D., Zavatone-Veth, J.A., and Clark, D.A. (2019). The manifold structure of limb coordination in walking *Drosophila*. eLife 8, e46409.
- Mathis, A., Mamidanna, P., Cury, K.M., Abe, T., Murthy, V.N., Mathis, M.W., and Bethge, M. (2018). DeepLabCut: markerless pose estimation of user-defined body parts with deep learning. Nat. Neurosci. 21, 1281–1289.
- Schnell, B., Weir, P.T., Roth, E., Fairhall, A.L., and Dickinson, M.H. (2014). Cellular mechanisms for integral feedback in visually guided behavior. Proc. Natl. Acad. Sci. USA 111, 5700–5705.
- Maisak, M.S., Haag, J., Ammer, G., Serbe, E., Meier, M., Leonhardt, A., Schilling, T., Bahl, A., Rubin, G.M., Nern, A., et al. (2014). A directional tuning map of Drosophila elementary motion detectors. Nature 500, 212–216.
- 11. Green, J., Vijayan, V., Mussells Pires, P., Adachi, A., and Maimon, G. (2019). A neural heading estimate is compared with an internal goal to guide oriented navigation. Nat. Neurosci. 22, 1460–1468.

Animal behaviour: Shifting attention in order to disperse

Laura Molina-García and Arantza Barrios*

Department of Cell and Developmental Biology, University College London, Rockefeller Building, 5th Floor, 21 University Street, London WC1E 6DE, UK *Correspondence: a.barrios@ucl.ac.uk

https://doi.org/10.1016/j.cub.2021.08.017

New findings in the nematode *Caenorhabditis elegans* identify neuromodulation of behavioural responses to pheromones as a mechanism for regulating dispersal and foraging strategies.

Dispersal as a foraging strategy (i.e. the search for new sources of food) is a behaviour critical for survival. Foraging in a constantly changing environment requires integration of information about food availability as well as social cues, such as pheromones, which signal the presence of potential competitors. The bacterivorous nematode *Caenorhabditis elegans* is a powerful system to study how the integration of food and pheromone signals shapes foraging behaviour.





Figure 1. Sexually dimorphic responses to the pheromone ascr#3 result in different foraging strategies. (A) Behavioural avoidance response of *C. elegans* hermaphrodites to the pheromone ascr#3 as well as its cellular (ASI neuron) and molecular (DAF-7) regulators. The left panel indicates a dampening of the avoidance response through PDF-1 modulation and according to food abundance, reducing dispersal from a foodrich environment. (B) Attractive behavioural response of *C. elegans* males to the pheromone ascr#3 (mediated by the ADF neuron); this response is not modified by food availability.

C. elegans has a 'boom and bust' life cycle in which food availability and population density change rapidly and dramatically¹. Therefore, worms need to collect and integrate information about these variables to decide whether to exploit a given environment or explore in search of a new one. Several factors have been shown to modulate C. elegans foraging and dispersal. These include food abundance^{2,3}, food nutritional value⁴, population density³ (which is sensed through blends of pheromones called ascarosides⁵), experience⁶ and biological sex. However, the neural mechanisms by which all these signals are integrated to modulate foraging remain poorly understood. In this issue of Current Biology, Luo and Portman⁷ provide new insight into the molecular and cellular regulation of foraging in a sexually dimorphic context (Figure 1).

By systematically changing the thickness of the bacterial lawn on which *C. elegans* feed, these authors find that responses to the pheromone ascr#3, which signals population density, are modulated by food abundance and in a

sex-specific manner. Hermaphrodites chronically avoid ascr#3, but avoidance is reduced when food is abundant. Males, instead, are attracted to ascr#38 and this response is independent of food abundance. Luo and Portman⁷ also show that integration of food signals and pheromones in hermaphrodites requires signalling by the neuropeptide pigment-dispersing factor (PDF). Mutant hermaphrodites lacking the PDF receptor PDFR-1 avoid ascr#3 even when plenty of food is available. Conversely, increasing PDFR-1 signalling, by ligand overexpression, reduces ascr#3 avoidance when food is limited⁷. Therefore, the intensity of PDFR-1 signalling, mainly in a PDF-1-dependent manner, could encode information about food abundance to modulate ascr#3 avoidance.

Where does PDFR-1 act to modulate ascr#3 avoidance? To answer this question, the authors used a previously described intersectional genetic strategy to restore PDFR-1 signalling in a cell-specific manner⁹. They show that PDFR-1 is required in the nervous system,

specifically in a small group of interneurons (AIA, PVC, AVA, AVD, AVE, AVG and RIM), likely by modulating the processing and perception of the pheromone signal rather than its sensation.

Next, the authors go on to investigate the neural substrates for ascr#3 detection. Ascr#3 is not only a population density marker but also a sex pheromone and, as mentioned earlier, males and hermaphrodites display very different responses to ascr#3: males show attraction, whereas hermaphrodites show repulsion^{8,10,11}. Such dimorphism may be explained by the need to incorporate reproduction demands into behavioural decisions. For self-fertilising hermaphrodites, whose reproductive priority is to find a suitable environment to lay eggs, high levels of pheromones indicate competition for resources. In contrast, for males, whose reproductive priority is to find a mating partner, pheromones indicate a potential source of mating opportunities.

The circuits mediating responses to ascr#3 are also sexually dimorphic.

Current Biology Dispatches

Previous studies showed that acute responses to ascr#3 in the absence of food require the ADL and ASK neurons in hermaphrodites and the ASK and CEM neurons in males^{8,10,11}. However, combining genetic ablation of neurons and behavioural analysis, Luo and Portman⁷ find that chronic avoidance of ascr#3 in hermaphrodites requires a different class of neurons, the ASI neurons. Regarding males, the Portman lab has previously shown that chronic attraction to ascr#3 is mediated by ADF neurons¹². In the present work, these authors further show that removing ASI function in males does not disrupt attraction to ascr#3. However, if ADF is removed in males, they now display repulsion to ascr#3, like hermaphrodites, and this is also mediated by ASI. Therefore, in males there is a latent circuit that mediates hermaphrodite-like responses to pheromones. Interestingly, a latent circuit driving pheromonedependent male-specific sexual behaviour has also been found in female mice¹³. A further interesting observation is that, although PDF signalling dampens ascr#3 avoidance in both hermaphrodites and ADF-ablated males, modulation of ascr#3 avoidance by food occurs only in hermaphrodites and not in males. This suggests that PDF signalling in males may be encoding something other than food abundance.

The TGFβ superfamily ligand DAF-7 is one of the modulators secreted by ASI, and, similarly to ASI-ablated animals, daf-7 mutant hermaphrodites exhibited no response to ascr#3 regardless of food thickness⁷. This suggests that DAF-7 may mediate the ASI-driven avoidance of ascr#3. Since previous studies have shown that pheromones as well as a scarcity of food downregulate the expression of DAF-7¹⁴, the studies of Luo and Portman⁷ suggest that low levels of DAF-7 may mediate dispersal away from a thin lawn of food also containing pheromone. This is somewhat in contrast to other work, which has shown that foraging and dispersal require high levels of DAF-7². Further work is needed to reconcile these findings; determination of when exactly DAF-7 is required to mediate dispersal in these different contexts may help. One possibility is that DAF-7 may play different roles during development and during adulthood to

regulate pheromone sensing and foraging. Indeed, functional ASI neurons and DAF-7 signalling are necessary during development in males so that proper pheromone responses are elicited during adulthood¹⁵.

One open question remaining from the work by Luo and Portman⁷ is what exactly does PDF signalling encode? The authors propose that PDF encodes information about food. This information may be relayed through mechanosensation of bacteria by ADE or detection of metabolic gases via RMG (both of these neurons express PDF-1)¹⁶. However, recent findings provide an alternative interpretation: Dal Bello et al.¹⁷ show that worms change their preference for pheromones from attraction to repulsion when food is scarce, and this switch in preferences is due to associative learning between pheromones and food. Therefore, PDF-mediated dampening of ascr#3 aversion in thick food lawns may be a consequence of learning and integration of pheromones with food. Indeed, a role for PDF-1 in mediating associative learning of salt with pheromones has been reported in males¹⁸. Furthermore, Luo and Portman⁷ show that PDF signalling modulates ascr#3 by acting on AIA, a class of interneurons previously shown to underlie the aversive associative learning of attractive stimuli with lack of food¹⁹. Another potential function of PDF proposed by Luo and Portman⁷ is the modulation of attention to sensory stimuli. We agree with this interpretation and would like to extend it by proposing that PDF may function to shift attention away from one salient stimulus towards another to reorganise the hierarchy of behavioural priorities. This proposed function for PDF is consistent with the findings of several previous studies reviewed in Flavell et al.²⁰ as well as with the current work by Luo and Portman⁷. When food is abundant, it may be adaptive for hermaphrodites to ignore population density (through PDF modulation) and instead exploit an enriched environment. However, when food is being depleted, pheromones may be more salient because population density indicates competition for resources. This model is tested by Luo and Portman⁷ with an elegant experiment in which they measure the dispersal rates from food

patches of different thickness (scarce or abundant) in the presence or absence of ascr#3. Indeed, they find that dispersal rates are highest when food is low and pheromone is high and that this requires ASI neurons.

Taken together, the work of Luo and Portman⁷ identifies sexually dimorphic neurogenetic mechanisms underlying the integration of food and social cues during foraging. The findings also open new and interesting questions about the role of neuromodulators and the underpinnings of their sexually dimorphic influence.

REFERENCES

- Félix, M.-A., and Duveau, F. (2012). Population dynamics and habitat sharing of natural populations of Caenorhabditis elegans and C. briggsae. BMC Biol. 10, 59.
- Milward, K., Busch, K.E., Murphy, R.J., de Bono, M., and Olofsson, B. (2011). Neuronal and molecular substrates for optimal foraging in Caenorhabditis elegans. Proc. Natl. Acad. Sci. USA 108, 20672–20677.
- Harvey, S.C. (2009). Non-dauer larval dispersal in Caenorhabditis elegans. J. Exp. Zool. B Mol. Dev. Evol. 312B, 224–230.
- Shtonda, B.B., and Avery, L. (2006). Dietary choice behavior in Caenorhabditis elegans. J. Exp. Biol. 209, 89–102.
- Ludewig, A.H., and Schroeder, F.C. (2013). Ascaroside signaling in C. elegans (January 18, 2013). In The C. elegans Research Community, WormBook, ed. (WormBook), https://doi.org/10.1895/wormbook.1.155.1
- Pradhan, S., Quilez, S., Homer, K., and Hendricks, M. (2019). Environmental programming of adult foraging behavior in C. elegans. Curr. Biol. 29, 2867–2879.e4.
- Luo, J., and Portman, D.S. (2021). Sexspecific, pdfr-1-dependent modulation of pheromone avoidance by food abundance enables flexibility in C. elegans foraging behavior. Curr. Biol. 31, 4449–4461.
- Srinivasan, J., Kaplan, F., Ajredini, R., Zachariah, C., Alborn, H.T., Teal, P.E.A., Malik, R.U., Edison, A.S., Sternberg, P.W., and Schroeder, F.C. (2008). A blend of small molecules regulates both mating and development in Caenorhabditis elegans. Nature 454, 1115–1118.
- Flavell, S.W., Pokala, N., Macosko, E.Z., Albrecht, D.R., Larsch, J., and Bargmann, C.I. (2013). Serotonin and the neuropeptide PDF initiate and extend opposing behavioral states in C. elegans. Cell 154, 1023–1035.
- Macosko, E.Z., Pokala, N., Feinberg, E.H., Chalasani, S.H., Butcher, R.A., Clardy, J., and Bargmann, C.I. (2009). A hub-and-spoke circuit drives pheromone attraction and social behaviour in C. elegans. Nature 458, 1171– 1175.



CellPress

- Jang, H., Kim, K., Neal, S.J., Macosko, E., Kim, D., Butcher, R.A., Zeiger, D.M., Bargmann, C.I., and Sengupta, P. (2012). Neuromodulatory state and sex specify alternative behaviors through antagonistic synaptic pathways in C. elegans. Neuron 75, 585–592.
- Fagan, K.A., Luo, J., Lagoy, R.C., Schroeder, F.C., Albrecht, D.R., and Portman, D.S. (2018). A single-neuron chemosensory switch determines the valence of a sexually dimorphic sensory behavior. Curr. Biol. 28, 902–914.e5.
- Kimchi, T., Xu, J., and Dulac, C. (2007). A functional circuit underlying male sexual behaviour in the female mouse brain. Nature 448, 1009–1014.

- Ren, P., Lim, C.S., Johnsen, R., Albert, P.S., Pilgrim, D., and Riddle, D.L. (1996). Control of C. elegans larval development by neuronal expression of a TGF-beta homolog. Science 274, 1389–1391.
- White, J.Q., and Jorgensen, E.M. (2012). Sensation in a single neuron pair represses male behavior in hermaphrodites. Neuron 75, 593–600.
- 16. Janssen, T., Husson, S.J., Meelkop, E., Temmerman, L., Lindemans, M., Verstraelen, K., Rademakers, S., Mertens, I., Nitabach, M., Jansen, G., and Schoofs, L. (2009). Discovery and characterization of a conserved pigment dispersing factor-like neuropeptide pathway in Caenorhabditis elegans. J. Neurochem. 111, 228–241.
- Dal Bello, M., Pérez-Escudero, A., Schroeder, F.C., and Gore, J. (2021). Inversion of pheromone preference optimizes foraging in C. elegans. eLife 10, e58144.

Current Biology

Dispatches

- Sammut, M., Cook, S.J., Nguyen, K.C.Q., Felton, T., Hall, D.H., Emmons, S.W., Poole, R.J., and Barrios, A. (2015). Glia-derived neurons are required for sex-specific learning in C. elegans. Nature 526, 385–390.
- Tomioka, M., Adachi, T., Suzuki, H., Kunitomo, H., Schafer, W.R., and Iino, Y. (2006). The insulin/PI 3-kinase pathway regulates salt chemotaxis learning in Caenorhabditis elegans. Neuron 51, 613–625.
- 20. Flavell, S.W., Raizen, D.M., and You, Y.-J. (2020). Behavioral states. Genetics 216, 315–332.

Neuroscience: Turbulent times for brain information processing

Giulio Bondanelli¹ and Stefano Panzeri^{1,2,*}

¹Neural Computation Laboratory, Center for Human Technologies, Istituto Italiano di Tecnologia, Genova, Italy ²Department of Neural Information Processing, Center for Molecular Neurobiology (ZMNH), University Medical Center Hamburg-Eppendorf (UKE), Hamburg, Germany

*Correspondence: stefano.panzeri@zmnh.uni-hamburg.de https://doi.org/10.1016/j.cub.2021.09.006

A recent study shows that rare long-range connections between brain areas may considerably improve transmission of information between areas. The study suggests that information may propagate better through long-range connections when neural activity exhibits turbulent dynamics.

Big whorls have little whorls Which feed on their velocity, And little whorls have lesser whorls And so on to viscosity.

Lewis F. Richardson, 1922

The brain is composed of billions of neurons interacting with each other through electrical impulses. The functions of the brain are made possible thanks to these complex interactions, which occur at multiple spatial scales, ranging from the cellular level to large brain networks¹. But how does the information about the external world get processed by networks at different spatial scales, and how do networks organized at different scales interact with one another? A new study by Deco *et al.*² published in this issue of *Current Biology* suggests that long-range connections between brain areas have a special role in facilitating the flow of information across neural circuits organized at different spatial scales when neural activity exhibits turbulent dynamics.

Physics has been highly successful at developing conceptual frameworks for understanding how natural interactions, or forces, between elements of a physical system determine its collective dynamics and properties. In the last fifty years such ideas from physics have trickled into neuroscience, providing a major conceptual framework for relating the structure of the brain to its dynamics and function. This has given neuroscientists important tools to study how brain functions emerge from the interactions between thousands of neurons. Paramount examples of this approach include Hopfield's attractor neural networks for memory storage and retrieval³, the study of how neural oscillations emerge from the couplings between excitatory and inhibitory neurons⁴ and of how interactions between areas of the cerebral cortex shape its global dynamics at rest^{5,6}.

Traditional theories from theoretical physics (e.g. thermodynamics) typically focus on the interactions between upclose elements, termed short-range interactions. Neglecting long-range interactions between elements that are far apart from each other is a convenient and extremely effective simplification for the description of a variety of physical phenomena⁷. However, long-range interactions are ubiquitous in nature and may become dominant in specific circumstances (e.g. the evolution of galaxies or the Great Red Spot of Jupiter can be explained using the physics of long-range systems').

